AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

1. (Currently Amended) Method of treating an underlying sause of a pathological condition characterized by an increased IL-1 and/or TNF-α level which comprises administering to a subject having such condition a therapeutically effective amount of at least one member selected from a first group consisting of discerein and rhein in a pharmaceutical dosage form;

wherein said pathological condition is an inflammation or autoimmune disease; and wherein said pathological condition is selected from the group consisting of psoriatic arthritis. Wegener's disease, granulomatosis, asthma, pulmonary emphysema, Paget's disease, bone metastases, atherosclerosis, myeloma and myeloid leukemia.

- 2. (Cancel)
- (Previously Presented) The method of claim 1 wherein said group member is diacerein.
- 5. (Previously Presented) The method of claim 1 wherein said therapeutically effective amount is a daily dose of about 25 to about 500 mg of said group member,
- 6. (Previously Presented) The method of claim 1 wherein each unit dose of said pharmaceutical dosage form contains about 5 to about 500 mg of said group member.
- 7. (Previously Presented) The method of claim 1 wherein each unit dose of said pharmaceutical dosage form contains about 20 mg to about 200 mg of said group member.
- (Previously Presented) The method of claim 1 wherein each unit dose of said pharmaceutical dosage form contains about 5 to about 100 mg of said group member.

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 (Previously Presented) The method of claim 1 where each unit dose of said pharmaceutical dosage form contains about 50 mg of said group member.

- (Previously Presented) The method of claim 1 wherein said dosage form is a capsule.
- 11. (Currently Amended) Method of treating an underlying eause of an inflammatory and/or autoimmune condition characterized by an increased IL-1 and/or TNF-a level which comprises modifying the production or action of proinflammatory cytokines including said IL-1 and/or TNF-a by administering to a subject having such condition a therapeutically effective amount of at least one member selected from a first group consisting of diacerein and rhein in a pharmaceutical dosage form.
- 12. (Previously Presented) The method of claim 11 wherein said group member is diacerein.
- (Previously Presented) The method of claim 11 wherein said condition is rheumatoid arthritis.
- 14. (Previously Presented) Method for reducing the synthesis of IL-1 and TNF-a which comprises administering to a subject a therapeutically effective amount of at least one member selected from the group consisting of diacerein and rhein in a pharmaceutical dosage form.
- 15. (Withdrawn) The method of claim 1, wherein the first group member is administered in association with a treatment to relieve symptomatic discomfort.
- 16. (Withdrawn) The method of claim 1, wherein said method of treatment further comprises also administering a therapeutically effective amount of at least one member selected from a second group consisting of: (i) an IL-1 synthesis inhibitor, (ii) a TNF- α synthesis inhibitor, (iii) a composition that lowers the levels of IL-1, (iv) a

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composition that lowers the level of TNF- α , (v) an IL-1 receptor antagonist, (vi) a TNF- α receptor antagonist, (vii) a composition that down-regulates the number of IL-1 receptors, (viii) a composition that down-regulates the number of TNF- α receptors, and (ix) a combination thereof.

- 17. (Withdrawn) The method of claim 16 wherein the first and second group members are co-administered.
- 18. (Withdrawn) The method of claim 16 wherein the first and second group members are administered in association with each other.
- 19. (Withdrawn) The method of claim 1, wherein the need for surgery to treat said pathological condition is avoided, delayed or reduced by administering the first group member.
- 20. (Withdrawn) The method of claim 1, wherein the first group member is administered in association with a treatment to avoid, delay or reduce the need for surgery to treat said pathological condition.
- 21. (Withdrawn) The method of claim 11 wherein the first group member is administered in association with the treatment to relieve symptomatic discomfort.
- 22. (Withdrawn) The method of claim 11, wherein the need for surgery to treat said pathological condition is avoided, delayed or reduced by administering the first group member.
- 23. (Withdrawn) The method of claim 11, wherein the first group member is administered in association with a treatment to avoid, delay or reduce the need for surgery to treat said pathological condition.